- (e) Claims information. A Part D sponsor must furnish directly to enrollees, in the manner specified by CMS and in a form easily understandable to such enrollees, a written explanation of benefits when prescription drug benefits are provided under qualified prescription drug coverage. The explanation of benefits must—
- (1) List the item or service for which payment was made and the amount of the payment for each item or service.
- (2) Include a notice of the individual's right to request an itemized statement.
- (3) Include the cumulative, year-to-date total amount of benefits provided, in relation to—
- (i) The deductible for the current year.
- (ii) The initial coverage limit for the current year.
- (iii) The annual out-of-pocket threshold for the current year.
- (4) Include the cumulative, year-todate total of incurred costs to the extent practicable.
- (5) Include any applicable formulary changes for which Part D plans are required to provide notice as described in §423.120(b)(5).
- (6) Be provided during any month when prescription drug benefits are provided under this part, including for covered Part D spending between the initial coverage limit described in §423.104(d)(3) and the out-of-pocket threshold described in §423.104(d)(5)(iii).

§ 423.132 Public disclosure of pharmaceutical prices for equivalent drugs.

- (a) General requirements. Except as provided under paragraph (c) of this section, a Part D sponsor must require a pharmacy that dispenses a covered Part D drug to inform an enrollee of any differential between the price of that drug and the price of the lowest priced generic version of that covered Part D drug that is therapeutically equivalent and bioequivalent and available at that pharmacy, unless the particular covered Part D drug being purchased is the lowest-priced therapeutically equivalent and bioequivalent version of that drug available at that pharmacy.
- (b) Timing of notice. Subject to paragraph (d) of this section, the informa-

- tion under paragraph (a) of this section must be provided after the drug is dispensed at the point of sale or, in the case of dispensing by mail order, at the time of delivery of the drug.
- (c) Waiver of public disclosure requirement. CMS waives the requirement under paragraph (a) of this section in the case of—
- (1) An MA private fee-for-service plan described in §422.4 of this chapter that—
- (i) Offers qualified prescription drug coverage and provides plan enrollees with access to covered Part D drugs dispensed at all pharmacies, without regard to whether they are contracted network pharmacies; and
- (ii) Does not charge additional costsharing for access to covered Part D drugs dispensed at out-of-network pharmacies.
 - (2) An out-of-network pharmacy;
 - (3) An I/T/U network pharmacy;
- (4) A network pharmacy that is located in any of the U.S. territories; and
- (5) Other circumstances where CMS deems compliance with the requirements of paragraph (a) of this section to be impossible or impracticable.
- (d) Modification of timing requirement. CMS modifies the requirement under paragraph (b) of this section as follows—
- (1) For long-term care network pharmacies, which must meet the requirement in paragraph (a) of this section by providing such information to Part D plans for inclusion in the written explanations of benefits required under § 423.128(e); and
- (2) Under other circumstances where CMS deems compliance with the requirement under paragraph (b) of this section to be impossible or impracticable.

§ 423.136 Privacy, confidentiality, and accuracy of enrollee records.

For any medical records or other health and enrollment information it maintains with respect to enrollees, a PDP sponsor must establish procedures to do the following—

(a) Abide by all Federal and State laws regarding confidentiality and disclosure of medical records, or other health and enrollment information. The PDP sponsor must safeguard the